

In the Claims

Please amend the claims as follows. Applicant submits a new complete claim set in which insertions and deletions in marked-up claims are indicated by underlining and strikeouts, respectively.

1-36. (Canceled)

37. (Currently amended) A method of stimulating or enhancing a protective immune response to an antigen in a mammal; which method comprises administering ~~co-administering~~ to a mucosal surface of said mammal with the antigen an effective adjuvant amount of a non-toxic double mutant form of pertussis toxin, said antigen being one which elicits a protective immune response when administered ~~co-administered~~ with said effective adjuvant amount of said non-toxic double mutant form of pertussis toxin, wherein said non-toxic double mutant form of pertussis toxin comprises an S₁ sub unit containing an amino acid at position 129 which is other than glutamic acid and containing an amino acid at position 9 which is other than arginine.

38. (Canceled)

39. (Previously presented) A method according to claim 37, wherein the amino acid at position 129 in the S₁ sub-unit is glycine.

40. (Canceled)

41. (Previously presented) A method according to claim 37 wherein the amino acid at position 9 is lysine.

42. (Previously presented) A method according to claim 37 wherein the antigen and the non-toxic form of pertussis toxin are administered intranasally.

43. (Currently amended) A method according to claim 37 wherein the antigen and the non-toxic double mutant form of pertussis toxin are administered simultaneously or sequentially ~~at the same time~~.

44. (Previously presented) A method according to claim 43 wherein the antigen and the non-toxic double mutant form of pertussis toxin are present in admixture in a composition administered to the mammal.
45. (Previously presented) A method according to claim 37 wherein the antigen is selected from the group consisting of tetanus toxin C-fragment, and one or more immunogenic fragments thereof.
46. (Previously presented) A method according to claim 37 wherein the antigen is selected from the group consisting of FHA and P69.
47. (Withdrawn) A vaccine composition in the form of nasal drops or a nasal spray, the composition comprising an antigen and an adjuvant capable of enhancing the immune response to the antigen in a mammal to which the composition is administered; wherein the adjuvant is a non-toxic double mutant form of pertussis toxin and said antigen is one which elicits a protective immune response when co-administered with said adjuvant.
48. (Withdrawn) A vaccine composition according to claim 47 wherein the mutant form of pertussis toxin comprises an S₁ sub unit containing an amino acid at position 129 which is other than glutamic acid.
49. (Withdrawn) A vaccine composition according to claim 48 wherein the amino acid at position 129 is glycine.
50. (Withdrawn) A vaccine composition according to claim 47 wherein the non-toxic double mutant form of pertussis toxin comprises an S₁ sub unit containing an amino acid at position 9 which is other than arginine.
51. (Withdrawn) A vaccine composition according to claim 50 wherein the amino acid at position 9 is lysine.
52. (Withdrawn) A vaccine composition according to claim 47 packaged in a container for dispensing a metered dose of the composition in spray or drop form.

53. (Withdrawn) A vaccine composition comprising an antigen and an adjuvant which enhances the immune response to the antigen in a mammal to which the composition is administered; wherein the adjuvant is a non-toxic double mutant form of pertussis toxin and the antigen is selected from tetanus toxin C fragment and one or more immunogenic fragments thereof.

54. (Withdrawn) A vaccine composition according to claim 53 which is in a form selected from nasal drops and a nasal spray.

55. (Currently amended) A method according to claim 46 wherein both FHA and P69 are administered ~~co-administered~~ with the non-toxic double mutant form of pertussis toxin.